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(54) Title: N-[(PIPERAZINYL)HETARYL]ARYLSULFONAMIDE COMPOUNDS WITH AFFINITY FOR THE DOPAMINE D3 RECEPTOR

$$R^{1}-N$$
 $N-Q-N-SO_{2}-Ar$ 
 $(I)$ 
 $R^{2}$ 

(57) Abstract: The invention relates to N-[(piper-azinyl)hetaryl]arylsulfonamide compounds of the general formula (I) in which Q is a bivalent, 6-membered heteroaromatic radical which possesses 1 or 2 N atoms as ring members and which optionally carries one or two substituents R<sup>a</sup> which is/are selected, independently of each other, from halogen, CN, NO<sub>2</sub>, CO<sub>2</sub>R<sup>4</sup>, COR<sup>5</sup>, C<sub>1</sub>-C<sub>4</sub>-alkyl and C<sub>1</sub>-C<sub>4</sub>-haloalkyl; Ar is phenyl or a

6-membered heteroaromatic radical which possesses 1 or 2 N atoms as ring members and which optionally carries one or two substituents R<sup>b</sup>, which is/are selected from halogen, NO<sub>2</sub>, CN, CO<sub>2</sub>R<sup>4</sup>, COR<sup>5</sup>, C<sub>1</sub>-C<sub>6</sub>-alkyl, C<sub>2</sub>-C<sub>6</sub>-alkenyl, C<sub>2</sub>-C<sub>6</sub>-alkynyl, C<sub>3</sub>-C<sub>6</sub>-cycloalkyl-C<sub>1</sub>-C<sub>4</sub>-alkyl and C<sub>1</sub>-C<sub>4</sub>-haloalkyl, with it also being possible for two radicals R<sup>b</sup> which are bonded to adjacent C atoms of Ar to be together C<sub>3</sub>-C<sub>4</sub>-alkylene; R<sup>1</sup> is hydrogen, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy-C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>3</sub>-C<sub>4</sub>-alkenyl or C<sub>3</sub>-C<sub>4</sub>-alkynyl; with the radicals n, R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> having the meanings given in the patent claims, to the N-oxides and to the physiologically tolerated acid addition salts of these compounds and to pharmaceutical compositions which comprise at least one N-[(piperazinyl)hetaryl]arylsulfonamide compound as claimed in one of claims 1 to 10 and/or at least one physiologically tolerated acid addition salt of I and/or an N-oxide of I, where appropriate together with physiologically acceptable carriers and/or auxiliary substances for treating diseases which respond to influencing by dopamine D<sub>3</sub> receptor antagonists or agonists, in particular for treating diseases of the central nervous system and disturbances of kidney function.

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